Claims

- 1. An antibody binding to 40-kDa OMP or a functional fragment thereof, which has activity of inhibiting the binding of hemin to 40-kDa OMP.
- 2. An antibody binding to 40-kDa OMP or a functional fragment thereof, which has (1) activity of inhibiting the coaggregation of *P. gingivalis* and (2) activity of promoting human neutrophilic phagocytosis.
- 3. An antibody binding to 40-kDa OMP or a functional fragment thereof, which has (1) activity of inhibiting the coaggregation of *P. gingivalis* and (2) activity of inhibiting the binding of hemin to 40-kDa OMP.
- 4. An antibody binding to 40-kDa OMP or a functional fragment thereof, which has (1) activity of promoting human neutrophilic phagocytosis and (2) activity of inhibiting the binding of hemin to 40-kDa OMP.
- 5. An antibody binding to 40-kDa OMP or a functional fragment thereof, which has (1) activity of inhibiting the coaggregation of *P. gingivalis*, (2) activity of promoting human neutrophilic phagocytosis, and (3) activity of inhibiting the binding of hemin to 40-kDa OMP.
- 6. The antibody or the functional fragment thereof according to any one of claims 2, 3, and 5, wherein the coaggregation of *P. gingivalis* is coaggregation of *P. gingivalis* and *Actinomyces viscosus*.
- 7. An antibody binding to 40-KDa OMP or a functional fragment thereof, which has activity of suppressing alveolar bone resorption.
- 8. The antibody or the functional fragment thereof according to any one of claims 1 to 7, wherein the antibody is a human antibody.
- 9. The antibody or the functional fragment thereof according to any one of claims 1 to 8, which is produced by a mouse-mouse hybridoma.
- 10. The antibody or the functional fragment thereof according to any one of claims 1 to 9, wherein the antibody is a monoclonal antibody.
- 11. The antibody or the functional fragment thereof according to any one of claims 1 to 10, which covalently or non-covalently binds to a therapeutic

agent.

- 12. The antibody or the functional fragment thereof according to claim 11, wherein the therapeutic agent is selected from antibiotics or antibacterial agents.
- 13. The antibody or the functional fragment thereof according to claim 12, wherein the antibiotic or the antibacterial agent is tetracycline or minocycline.
- 14. The antibody or the functional fragment thereof according to any one of claims 1 to 13, wherein the antibody class is IgG.
- 15. The antibody or the functional fragment thereof according to claim 14, wherein IgG is IgG1.
- 16. The antibody or the functional fragment thereof according to any one of claims 1 to 13, wherein the antibody class is IgA.
- 17. The antibody or the functional fragment thereof according to any one of claims 1 to 16, wherein the amino acid sequence of a heavy chain constant region is altered.
- 18. An antibody binding to 40-kDa OMP or a functional fragment thereof, which is produced by a hybridoma h13-17 (accession No. FERM BP-8325).
- 19. An antibody binding to 40-kDa OMP or a functional fragment thereof, which comprises variable regions of an antibody that is produced by a hybridoma h13-17 (accession No. FERM BP-8325).
- 20. The antibody or the functional fragment thereof according to claim 18 or 19, which covalently or non-covalently binds to a therapeutic agent.
- 21. The antibody or the functional fragment thereof according to claim 20, wherein the therapeutic agent is selected from antibiotics or antibacterial agents.
- 22. The antibody or the functional fragment thereof according to claim 21, wherein the antibiotic or the antibacterial agent is tetracycline or minocycline.

- 23. The antibody or the functional fragment thereof according to any one of claims 18 to 22, wherein the antibody class is IgG.
- 24. The antibody or the functional fragment thereof according to claim 23, wherein IgG is IgG1.
- 25. The antibody or the functional fragment thereof according to any one of claims 18 to 22, wherein the antibody class is IgA.
- 26. The antibody or the functional fragment thereof according to any one of claims 18 to 25, wherein the amino acid sequence of a heavy chain constant region is altered.
- 27. A hybridoma h13-17 (accession No. FERM BP-8325).
- 28. An antibody binding to 40-kDa OMP or a functional fragment thereof, which is produced by a hybridoma 5-89-2 (accession No. FERM BP-8323).
- 29. An antibody binding to 40-kDa OMP or a functional fragment thereof, which comprises variable regions of an antibody that is produced by a hybridoma 5-89-2 (accession No. FERM BP-8323).
- 30. The antibody or the functional fragment thereof according to claim 28 or 29, which covalently or non-covalently binds to a therapeutic agent.
- 31. The antibody or the functional fragment thereof according to claim 30, wherein the therapeutic agent is selected from antibiotics or antibacterial agents.
- 32. The antibody or the functional fragment thereof according to claim 31, wherein the antibiotic or the antibacterial agent is tetracycline or minocycline.
- 33. The antibody or the functional fragment thereof according to any one of claims 28 to 32, wherein the antibody class is IgG.
- 34. The antibody or the functional fragment thereof according to claim 33, wherein IgG is IgG1.
- 35. The antibody or the functional fragment thereof according to any one of claims 28 to 32, wherein the antibody class is IgA.

- 36. The antibody or the functional fragment thereof according to any one of claims 28 to 35, wherein the amino acid sequence of a heavy chain constant region is altered.
- 37. A hybridoma 5-89-2 (accession No. FERM BP-8323).
- 38. An antibody binding to 40-kDa OMP or a functional fragment thereof, which is produced by a hybridoma a44-1 (accession No. FERM BP-8324).
- 39. An antibody binding to 40-kDa OMP or a functional fragment thereof, which comprises variable regions of an antibody that is produced by a hybridoma a44-1 (accession No. FERM BP-8324);
- 40. The antibody or the functional fragment thereof according to claim 40 or
- 41, which covalently or non-covalently binds to a therapeutic agent.
- 41. The antibody or the functional fragment thereof according to claim 40, wherein the therapeutic agent is antibiotics or antibacterial agents.
- 42. The antibody or the functional fragment thereof according to claim 41, wherein the antibiotic or the antibacterial agent is tetracycline or minocycline.
- 43. The antibody or the functional fragment thereof according to any one of claims 38 to 42, wherein the antibody class is IgG.
- 44. The antibody or the functional fragment thereof according to claim 43, wherein IgG is IgG1.
- 45. The antibody or the functional fragment thereof according to any one of claims 38 to 42, wherein the antibody class is IgA.
- 46. The antibody or the functional fragment thereof according to any one of claims 38 to 45, wherein the amino acid sequence of a heavy chain constant region is altered.
- 47. A hybridoma a44-1 (accession No. FERM BP-8324).
- 48. A nucleic acid, which is possessed by a hybridoma selected from the group consisting of a hybridoma h13-17 (accession No. FERM BP-8325), a hybridoma 5-89-2 (accession No. FERM BP-8323), and a hybridoma a44-1

(accession No. FERM BP-8324) and encodes an antibody containing a variable region of an antibody produced by the hybridoma or a functional fragment of the said antibody.

- 49. A protein encoded by the nucleic acid according to claim 48, which is an antibody or a functional fragment thereof.
- 50. An expression vector, which has the nucleic acid according to claim 48.
- 51. A host, which has the expression vector according to claim 50.
- 52. The host according to claim 51, which is selected from the group consisting of *Escherichia coli*, yeast cells, insect cells, mammalian cells, plant cells, and mammals.
- 53. A method for producing an antibody binding to 40-kDa OMP, which comprises isolating a gene that encodes an antibody binding to 40-kDa OMP from a hybridoma selected from the group consisting of a hybridoma h13-17 (accession No. FERM BP-8325), a hybridoma 5-89-2 (accession No. FERM BP-8323), and a hybridoma a44-1 (accession No. FERM BP-8324), constructing an expression vector comprising the gene, introducing the expression vector into a host to cause expression of the antibody, and collecting the antibody from the obtained host, the culture supernatant of the host, or secretion from the host.
- 54 An agent for suppressing alveolar bone resorption, which contains an antibody binding to 40-KDa OMP or a functional fragment thereof as an active ingredient.
- 55. An agent for preventing, diagnosing, or treating periodontal diseases, which contains an antibody binding to 40-kDa OMP or a functional fragment thereof as an active ingredient.
- 56. Use of an antibody binding to 40-KDa OMP or a functional fragment thereof for production of an agent for suppressing alveolar bone resorption.
- 57. A method for suppressing alveolar bone resorption, which comprises preparing an antibody binding to 40-KDa OMP or a functional fragment

thereof and administering the antibody or the fragment to an animal.

- 58. Use of an antibody binding to 40-KDa OMP or a functional fragment thereof for production of an agent for preventing, diagnosing, or treating periodontal diseases.
- 59. A method for diagnosing, preventing, or treating periodontal diseases, which comprises preparing an antibody binding to 40-KDa OMP or a functional fragment thereof and administering the antibody or the fragment to an animal.
- 60. An agent for preventing, diagnosing, or treating periodontal diseases, which contains the antibody or the functional fragment thereof according to any one of claims 1 to 26, 28 to 36, 38 to 46, and 49 as an active ingredient.
- 61. An agent for suppressing alveolar bone resorption, which contains the antibody or the functional fragment thereof according to any one of claims 1 to 26, 28 to 36, 38 to 46, and 49 as an active ingredient.
- 62. Use of the antibody or the functional fragment thereof according to any one of claims 1 to 26, 28 to 36, 38 to 46, and 49 for production of an agent for preventing, diagnosing, or treating periodontal diseases;
- 63. Use of the antibody or the functional fragment thereof according to any one of claims 1 to 26, 28 to 36, 38 to 46, and 49 for production of an agent for suppressing alveolar bone resorption.
- 64. A method for diagnosing, preventing, or treating periodontal diseases, which comprises preparing the antibody or the functional fragment thereof according to any one of claims 1 to 26, 28 to 36, 38 to 46, and 49 and administering the antibody or the fragment to an animal.
- 65. A method for suppressing alveolar bone resorption, which comprises preparing the antibody or the functional fragment thereof according to any one of claims 1 to 26, 28 to 36, 38 to 46, and 49 and administering the antibody or the fragment to an animal.